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Effect of dietary lipid structure in early postnatal life on mouse adipose tissue development and function in adulthood

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Abstract

Obese individuals have more (hyperplastic) and larger (hypertrophic) adipocytes in their white adipose tissue (WAT) than normal-weight individuals. The difference in cell number emerges early in childhood, suggesting that this is a critical period for being susceptible to obesity. Breast-feeding has been shown to be protective against obesity, and we have previously shown in mice that the physical structure of lipids in human milk may contribute to this protective effect. In the present study, we investigated how differences in the physical structure of lipids in the early diet may modulate adipose tissue development. Male mice were fed a diet containing control infant milk formula (Control IMF; Danone Research) or Nuturis[®] (Concept IMF with large phospholipid-coated lipid droplets; Danone Research) from post-natal day (PN)16 to 42. Subsequently, mice were challenged with a moderate Western-style diet (WSD) until PN98, and body composition was monitored by dual-energy X-ray absorptiometry. Epididymal WAT was analysed for adipocyte size, number and gene expression of metabolic transcription factors. Early Concept IMF exposure reduced fat accumulation during the WSD challenge by 30% compared with the Control IMF. It reduced adipocyte size without affecting adipocyte number in adult mice. The Concept IMF decreased the expression of PPAR γ , CCAAT/enhancer-binding protein and retinoid X receptor α in WAT in adulthood, key regulators of metabolic activity. In conclusion, Concept IMF exposure in early life reduced susceptibility to obesity in adult life, by preventing adipocyte hypertrophy upon adult dietary challenge without affecting adipogenesis. These data emphasise the importance of the physical properties of dietary lipids in early life in obesity risk later in life.

Key words: Nutritional programming: Obesity prevention: Dietary lipids: Infant nutrition

The development of human white adipose tissue (WAT) starts in the third trimester of gestation and continues in postnatal life through the proliferation of preadipocytes and subsequent differentiation, resulting in mature adipocytes storing energy in the form of lipids⁽¹⁾. Adipocyte number increases throughout childhood and adolescence to plateau and remain constant in adulthood^(2,3). Surprisingly, adult cell turnover is similar in obese and lean subjects, although adult obese subjects have more adipocytes compared with lean subjects. Interestingly, this difference in adipocyte number already emerges early in childhood, suggesting that this may be a critical period for WAT development and obesity risk^(2,3).

Comparable to human ontogeny, WAT development in rodents is initially driven by hyperplastic growth (increase in number) and subsequent hypertrophic growth (increase in cell size). When fed a standard chow, total cell number is established in rodents between 9 and 18 weeks of age^(4–6), whereas adipocyte size can increase until senescence⁽⁷⁾. Additionally, adipocyte size and number are higher in obese rats compared with lean rats⁽⁸⁾, and hyperplastic and hypertrophic growth dynamics differ according to fat depot, sex and diet^(9,10).

The growth of both hypertrophic and hyperplastic adipocytes is regulated by the concerted actions of several

Abbreviations: %FM, percentage of fat mass; BW, body weight; CEBP, CCAAT/enhancer-binding protein; EPI, epididymal; FA, fatty acid; FM, fat mass; HDL-C, HDL-cholesterol; IMF, infant milk formula; ING, inguinal; LDL-C, LDL-cholesterol; PL, phospholipids; PN, postnatal day; RXR α , retinoid X receptor α ; SREBP1c, sterol regulatory element-binding protein 1c; TC, total cholesterol; WAT, white adipose tissue; WSD, Western-style diet.

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transcription factors such as PPAR γ , retinoid X receptor α (RXR α) and the CCAAT/enhancer-binding protein (CEBP) family members^(11,12). *In vitro* studies have shown that PPAR γ -deficient embryonic stem cells are unable to differentiate into adipocytes⁽¹³⁾. Additionally, overexpression of PPAR γ in mature 3T3-L1 adipocytes reduces lipolysis and increases cell size and TAG content compared with control adipocytes. In contrast, a dominant-negative PPAR γ mutant has the opposite effect of increasing lipolysis and reducing fatty acid (FA) uptake⁽¹⁴⁾. Data from human and mouse genetic studies have shown that the deletion of the WAT-specific PPAR γ 2 isomer reduces adipogenesis and impairs insulin sensitivity. Thus, PPAR γ is clearly a key regulator to maintain lipid metabolic homeostasis. Many PUFA and lipid-derived mediators can act as endogenous PPAR γ ligands⁽¹⁵⁾. This enables sensing of nutritional signals that can be translated into a metabolic response to maintain homeostasis.

Experimental studies have provided compelling evidence for the sustained effects of early-life nutrition on adipose tissue development and function. For instance, both maternal obesity and intra-uterine growth retardation induced by a 50% food-restricted diet in rats resulted in the up-regulation of PPAR γ , RXR α and CEBP family members in the offspring directly after birth, which was associated with increased adiposity during adulthood^(16,17). A clinical study investigating the obesogenic effect of a specific human PPAR γ polymorphism has shown that breast-feeding is correlated with a lower BMI in adolescents carrying this polymorphism, suggesting that the protective effects of breast-feeding on the development of obesity are mediated, at least in part, via PPAR γ ⁽¹⁸⁾.

The protective effects of breast-feeding on obesity and metabolic health later in life are probably multifactorial⁽¹⁹⁾. One specific aspect is the fact that human milk contains large lipid droplets with an average mode diameter of 4 μ m surrounded by a native biological membrane composed of phospholipids (PL), proteins, enzymes, cholesterol and glycoproteins⁽²⁰⁾. In contrast, infant milk formula (IMF) contains small lipid droplets with an average mode diameter of 0.3–0.5 μ m lacking such a biological membrane coating. We have previously shown that this physical lipid structure may play a role in the programming of obesity risk later in life⁽²¹⁾. We developed a concept IMF in which the lipid droplets had physical properties similar to those of lipid droplets in human milk, i.e. larger in size and coated with PL (Nuturis[®]; Danone Research), and investigated whether postnatal exposure to this Concept IMF protects against excessive fat accumulation in adulthood. Although adult body weight (BW) was comparable between the Control IMF- and Concept IMF-fed mice challenged by a mild Western-style diet (WSD) from adolescence, adult fat mass (FM) was 28% lower in the mice fed the Concept IMF in early life. This was accompanied by an improved metabolic profile since adult fasting plasma lipids as well as plasma glucose, resistin and leptin levels were lower in the Concept IMF group. Taken together, these data suggest that the sustained effects on adult FM were mediated through differences in WAT development. We hypothesised that the exposure of pre-weaning mice to the Concept IMF with large PL-coated lipid droplets alters WAT

development by limiting hyperplastic and hypertrophic growth affecting the metabolic functions of WAT and ultimately leading to reduced adult body fat accumulation.

In the present study, we investigated the morphological WAT characteristics and gene expression of transcription factors that play a key role in adipocyte differentiation and lipogenesis to gain more insights into the mechanism underlying the programming effects of the Concept IMF on adult body composition.

Methods

Animals and study design

All experimental procedures were approved by the Animal Experimental Committee DEC-Consult, Bilthoven, The Netherlands, and complied with the principles of laboratory animal care. Mice were housed at Wageningen University and Research Centre (CKP, WUR, Wageningen, The Netherlands) under a 12 h light–12 h dark cycle (lights on at 06.00 hours = Zeitgeber time 0 h) in a temperature- and humidity-controlled room (21 \pm 2°C and 50 \pm 5%, respectively). Food and water were available *ad libitum* during the entire experimental protocol. BW was measured per litter before weaning and individually after weaning twice a week. Male and female C57BL/6J mice were time-mated. Dams were assigned to the American Institute of Nutrition-93G diet during pregnancy and lactation. After birth, pups were randomly cross-fostered, i.e. all pups from one litter were assigned to different dams, and litters were culled to four male and two female pups on postnatal day (PN)2. Litters were randomly assigned to one of the two experimental diets on PN16. After weaning on PN21, female pups were killed; male pups were housed in pairs and continued to be fed their respective diets until PN42, followed by a WSD. Food intake was measured weekly per cage between PN42 and 98.

To address the effects of the WSD challenge in adolescence and adulthood, we included a separate non-challenged reference group (Reference), which was maintained on the Control IMF until PN42 and then fed a standard semi-synthetic rodent chow until dissection on PN98 (Fig. 1). BW, FM and lean body mass were measured on PN42, 70 and 98 by dual-energy X-ray absorptiometry (PIXImus; GE Lunar) under general anaesthesia (isoflurane/N₂O/O₂). Relative FM was calculated as the percentage of FM (%FM) of total BW.

Infant milk formula production and analyses

As a basis for the Control IMF rodent diet, an IMF powder was prepared according to a standard stage 1 IMF recipe (for infants aged between 0 and 6 months) and a processing procedure (Danone Research). The Concept IMF powder that served as a basis for the Concept IMF rodent diet was prepared by adding 0.5 g/l PL of dairy origin (Fonterra Co-operative Group Limited; Table 1) and applying an altered processing procedure to obtain PL-coated lipid droplets larger than those in a standard IMF. Briefly, the polar PL were added to the aqueous phase and blended with the lipid phase containing the neutral vegetable lipids to generate a PL coating around the lipid droplets. Homogenisation

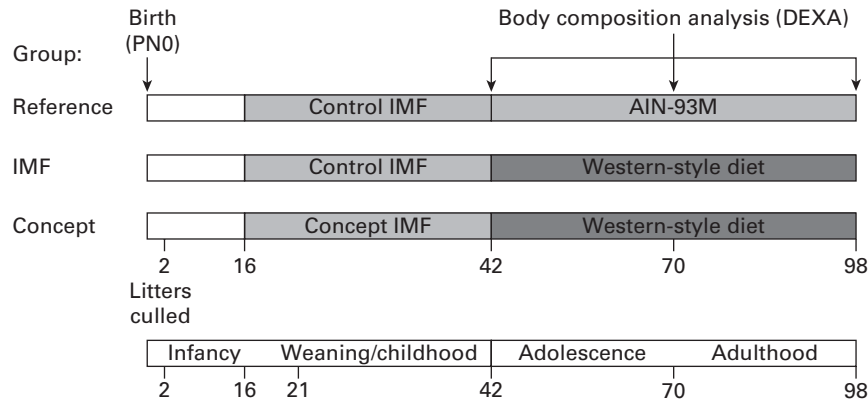


Fig. 1. Study design. DEXA, dual-energy X-ray absorptiometry; IMF, infant milk formula; PN, postnatal day.

pressure during processing was adjusted to obtain a homogeneous mixture of ingredients and also to retain a large lipid droplet size⁽²²⁾. Initial differences in FA composition between diets as a result of the dairy origin of the PL source that were not corrected for in the previous study⁽²²⁾ were compensated for by minor adjustments to the composition of the vegetable oil blend. Both IMF recipes contained 0.35 g arachidonic acid and 0.2 g DHA/100 g total FA.

Lipid droplet size distribution of the IMF powders was measured using a Mastersizer laser light-scattering analyser with a refractive index ratio of 1.46 (Mastersizer 2000, Hydro 2000G, Malvern Instruments Limited). PL in the coated lipids were visualised using epifluorescence microscopy (Carl Zeiss Benelux). Solubilised IMF was incubated with Annexin-V Alexa Fluor[®] 488 (Invitrogen – Molecular Probes) and Nile Red (Sigma-Aldrich) according to the manufacturer's protocol for staining PL and TAG, respectively, and labelling of the fat core and PL coating in the non-fixed lipid droplets. Analyses were performed using a fluorescence microscope (Zeiss Axioplan 2; Carl Zeiss) with a (10) × 63 oil immersion objective using fluorescein isothiocyanate (FITC) and tetramethylrhodamine isothiocyanate (TRITC) optical filters (Carl Zeiss Benelux).

Experimental diets

All experimental diets were semi-synthetic (Research Diet Services; Table 1) and had a macronutrient and micronutrient composition according to the American Institute of Nutrition formulation of American Institute of Nutrition-93G purified diets for laboratory rodents⁽²³⁾. The postnatal IMF-based rodent diets contained 28.3 w/w% Control or Concept IMF powder, respectively, and were complemented with protein and carbohydrates to meet rodent nutrient requirements. These diets were not pelletised, but given to the mice as dough (powder mixed with water) in order to preserve the lipid structure. The moderate WSD consisted of 10% (w/w) fat (5% (w/w) lard and 0.1% (w/w) cholesterol).

Blood sampling and dissection

Blood was sampled by cheek puncture under anaesthesia (isoflurane/N₂O/O₂) on PN42 after 4 h of fasting during the

light phase⁽²⁴⁾. Plasma was pooled per three mice per group to obtain sufficient samples for analyses. At dissection (PN98), mice were terminally anaesthetised (isoflurane/N₂O/O₂) after 4 h of fasting. Dual-energy X-ray absorptiometry scans were

Table 1. Composition of the experimental diets

	IMF			
	Control	Concept	WSD	AIN-93M
Carbohydrates (g/kg)	645	645	600	720
Sugars (g/kg)	235	235	150	100
Dextrose (g/kg)	5	5	150	–
Lactose (g/kg)	145	145	–	–
Sucrose (g/kg)	85	85	–	100
Polysaccharides (g/kg)	410	410	450	620
Maltodextrin (g/kg)	50	50	250	155
Starch (g/kg)	360	360	200	465
Protein (g/kg)	179	179	179	126
Soya (g/kg)	150	150	179	126
Whey (g/kg)	17	17	–	–
Casein (g/kg)	12	12	–	–
Fat (g/kg)	70	70	100	40
Oil blend* (g/kg)	69.1	64.4	50	–
Milk fat (g/kg)	0.9	5.6	–	–
Lard (g/kg)	–	–	50	–
Soya oil (g/kg)	–	–	–	40
SFA (g/kg)	28.7	29.7	41.9	5.9
MUFA (g/kg)	26.5	26.1	42.3	10.3
PUFA (g/kg)	11.5	11.0	13.2	23.5
LA (18:2n-6) (g/kg)	9.8	9.3	11.9	20.4
ALA (18:3n-3) (g/kg)	1.8	1.7	1.3	2.7
ARA (20:4n-6) (g/kg)	0.25	0.25	–	–
EPA (20:5n-3) (g/kg)	0.03	0.04	–	–
DHA (22:6n-3) (g/kg)	0.14	0.14	–	–
LA:ALA	5.4	5.3	9.5	7.5
Total n-6:n-3	5.1	5.0	9.5	7.5
Phospholipids† (g/kg)	0.1	1.1	–	–
Cholesterol (mg/kg)	4.7	20.4	1000	–
Cellulose	50	50	50	50
(Vitacel L 600-20) (g/kg)	–	–	–	–
Mineral mix (g/kg)	35	35	35	35
Vitamin mix (g/kg)	10	10	10	10
Choline bitartrate (g/kg)	2.5	2.5	2.5	2.5
L-Cysteine (g/kg)	3	3	3	3
tert-Butylhydroquinone (g/kg)	0.014	0.014	0.014	0.014

IMF, infant milk formula; WSD, Western-style diet; AIN, American Institute of Nutrition; LA, linoleic acid; ALA, α-linolenic acid; ARA, arachidonic acid.

* Mixture of oils, among others rapeseed, sunflower, fish, coconut and palm oils.

† Phospholipids derived from bovine milk.



performed, and blood was collected in K₃EDTA-coated tubes via cardiac puncture. Liver, pancreas, *m. tibialis*, brain and WAT depots (epididymal (EPI), retroperitoneal, perirenal and inguinal (ING)) were collected and weighed. Plasma was obtained by centrifugation at 1350 *g* for 12 min at 4°C and stored at −80°C.

Plasma analyses

Total cholesterol (TC), HDL-cholesterol (HDL-C), LDL-cholesterol (LDL-C), VLDL-cholesterol and TAG concentrations were determined colorimetrically after enzymatic conversion (Reinier de Graaf Laboratory).

Plasma glucose concentration (GOD-PAP method; Roche Diagnostics) was measured colorimetrically in PN98 plasma samples and analysed using a microplate imaging system (Bio-Rad Laboratories, Inc.). Plasma insulin concentration was measured by means of ELISA (DRG) according to the manufacturer's protocol. Homeostasis model assessment of insulin resistance was calculated from fasting plasma glucose and insulin concentrations (glucose (mmol/l) × insulin (pmol/l)/22.5) as an indirect measure of insulin sensitivity. Plasma volumes were insufficient to perform glucose and insulin analyses on PN42 samples.

Cytological and biochemical analyses of adipose tissue

Cell size distribution according to the optical method of Di Girolamo & Fine⁽²⁵⁾ was determined in fresh ING and EPI WAT depots, respectively, representing subcutaneous and abdominal fat. Depots were cut into large pieces of 1 mm and incubated in gassed (95% O₂/5% CO₂) Krebs–Ringer bicarbonate buffer (pH 7.4, 2 mg collagenase type II/ml; Gibco for Invitrogen) at 37°C. The digested tissue was filtered first through a 250 µm cell strainer, washed three times with Krebs–Ringer bicarbonate buffer and filtered again through a 40 µm cell strainer.

Images of isolated adipocytes were captured using an Axio-plan 2 Zeiss microscope (Carl Zeiss) and a Sony DXC-950P video camera (Sony). For the assessment of mean adipocyte volume and size distribution per mouse, six representative sections per slide and six slides per depot per mouse were used to obtain images of at least 300 cells using analysis software (Soft Imaging System).

The remaining WAT depots were snap-frozen and stored at −80°C for lipid content measurement. Lipid extraction was performed according to the method of Folch *et al.*⁽²⁶⁾ using dichloromethane–methanol (2:1, v/v).

WAT cellularity was calculated by dividing the fat pad lipid content by mean adipocyte weight. The mean adipocyte weight was calculated by multiplying the mean adipocyte volume by lipid density, which was derived from a triolein density (ρ) of 0.915 g/ml⁽²⁵⁾.

Gene expression analyses in epididymal white adipose tissue depot

Total RNA was isolated from epididymal WAT using the RNeasy Mini Kit (Qiagen). Complementary DNA (cDNA) was prepared

using the iScript cDNA Synthesis Kit (Bio-Rad), and quantitative real-time PCR analysis was performed using the CFX96 Real-Time PCR Detection System (Bio-Rad) with iQ SYBR Green Supermix (Bio-Rad). Validated quantitative PCR primers (Table 2) were obtained for *leptin* from SABiosciences (Qiagen) and for *PPAR* γ , *CEBP* α , *CEBP* β , *CEBP* δ , *RXR* α , sterol regulatory element-binding protein 1c (*SREBP1C*) and ribosomal protein S13 (*RPS13*) from Isogen Life Science. mRNA expression was calculated using the CFX Manager software (version 1.6; Biorad) and corrected for the expression of the housekeeping gene *RPS13*⁽²⁷⁾.

Statistical analyses

Statistical analyses were performed using SPSS 15.0.1 software (SPSS Benelux). Variables were checked for Gaussian distribution with the Shapiro–Wilk test. Levene's test for equality of variance was used to estimate the probability that the treatment groups had different variances. Direct and programming effects of any experimental diet (Control IMF, Concept IMF or Reference) on the development of body composition were analysed using a repeated-measures ANOVA. *Post hoc* analyses were performed on significant diet × time interactions using univariate ANOVA on PN42, 70 and 98 separately. Repeated-measures ANOVA was performed to analyse diet effects on the frequency distribution of adipocyte size on PN98 and a *post hoc* analysis was performed when significant diet × time interactions were confirmed. The effects of experimental diets on WAT weight, organ weight, plasma parameters, average adipocyte number and gene expression on PN98 were analysed by means of a univariate ANOVA.

Data are presented as means with their standard errors, unless otherwise indicated. Differences were considered significant at $P < 0.05$.

Results

Infant milk formula-based diets

The adjusted IMF processing and addition of 0.5 g/l PL from dairy origin resulted in a Concept IMF powder with PL-coated (Fig. 2) lipid droplets larger than those in the Control IMF (Table 3).

Direct effect of the Control and Concept IMF

BW gain in the Control IMF- and Concept IMF-fed mice was similar from PN2 to 42 ($F_{(2,31)} = 0.758$, $P = 0.477$; data not shown). Additionally, there was no direct effect of the diet during early development on body composition, since BW, lean body mass, FM and %FM were comparable (Fig. 3) on PN42. Plasma TAG, TC, HDL-C, LDL-C and VLDL-cholesterol concentrations were similar in all the groups (Table 5).

Programming effect of the Concept IMF during adult Western-style diet challenge

Body composition and food intake. BW gain during the WSD challenge was significantly lower in the Concept

Table 2. Quantitative PCR primers

Amplified region	Forward primer	Reverse primer	Annealing temperature (°C)
<i>PPARγ</i>	5'-AGATTAATAGTTTGACGGGGTTT-3'	5'-ACCCGTATCCCCTCTAATCGA-3'	62
<i>CEBPδ</i>	5'-AGAGCGCCATCGACTTCAGC-3'	5'-CCAAGCTCACCAGTGTCTGC-3'	63
<i>CEBPβ</i>	5'-TACCCAGGACCCATTGGATA-3'	5'-TTCACTTGGCCACTCTTCCT-3'	62
<i>CEBPα</i>	5'-TGTTGGAGTTGACCAAGTGAC-3'	5'-ATCCAGCGACCCCTAAACCAT-3'	60
<i>RXRα</i>	5'-GATGGCCTGTGTGGATCTTT-3'	5'-AACCAGCAACCAGAACCAAGC-3'	60
<i>SREBP1C</i>	5'-CAAGTGCTGCAGGAACTGA-3'	5'-CATGGCCTTGTCATGGAAC-3'	60
Leptin	SABiosciences®		63

CEBP, CCAAT/enhancer-binding protein; *RXR*, retinoid X receptor; *SREBP1C*, sterol regulatory element-binding protein 1c.

IMF mice compared with the Control IMF mice ($P=0.028$; Fig. 3). The BW difference was exclusively due to a lower FM gain in the Concept IMF mice compared with the Control IMF mice, since lean body mass gain from PN42 to 98 was similar in both the groups. Early postnatal Concept IMF exposure reduced FM gain due to the WSD challenge by 30% and %FM gain by 20% compared with Control IMF exposure. The %FM in the Concept IMF mice was similar to that in the non-challenged Reference mice ($P=0.234$) on PN70 and showed intermediate values on PN98 ($P=0.001$ and $P=0.012$) for the Control and non-challenged Reference mice, respectively (Fig. 4). The weights of WAT depots followed a similar pattern as total FM and %FM, with 26–34% lower WAT depot weights being observed in the Concept IMF mice compared with the Control IMF mice. Average liver weight of the Concept IMF mice was significantly lower than liver weight of the Control IMF mice and comparable to that of the Reference mice ($F(2,31) = 5.656$, $P=0.008$; Table 4). Food intake was similar in both the groups (53.09 (SEM 1.3) *v.* 50.88 (SEM 0.8) kJ/d per mouse for the Control and Concept IMF groups, respectively).

Adipocyte size and number. We aimed to address whether the differences in FM were related to a decrease in adipocyte number, adipocyte size or both in fat depots representing subcutaneous and abdominal fat^(9,10). The data on cell number

(Fig. 4) showed that this parameter was not affected by early dietary exposure; there was no difference in adipocyte number in the EPI and ING WAT depots of the Control IMF and Concept IMF groups.

The early diet did affect adipocyte size. Cell size distribution curves illustrate that mice fed the Concept IMF before the WSD challenge had a significantly smaller number of large adipocytes in the ING depot compared with the Control IMF mice ($F(14,266) = 3.199$, $P=0.002$). This difference was mainly found in the cell size category between 80 and 150 μm diameter. Average ING adipocyte volume was 25% lower in the Concept IMF group compared with the Control IMF group ($t(18) = 2.016$, $P<0.05$). Average EPI adipocyte volume at the end of the WSD challenge was 21% lower in the Concept IMF mice compared with the Control IMF mice, but this difference did not reach significance. As stated earlier, hyperplastic and hypertrophic growth of WAT is regulated by transcription factors, including *PPAR γ* , *RXR α* , *CEBP* and *SREBP1c*. The expression of *PPAR γ* ($F(1,7) = 4.701$, $P<0.01$), *RXR α* ($F(1,7) = 1.919$, $P<0.05$) and *CEBP α* ($F(1,3) = 15.90$, $P<0.001$) was decreased in the EPI WAT depots of the Concept IMF compared with the Control IMF mice. No differences in the expression of *CEBP β* , *CEBP δ* or *SREBP1c* were observed between both the IMF groups. The expression of leptin was significantly lower ($F(1,7) = 2.077$, $P<0.05$)

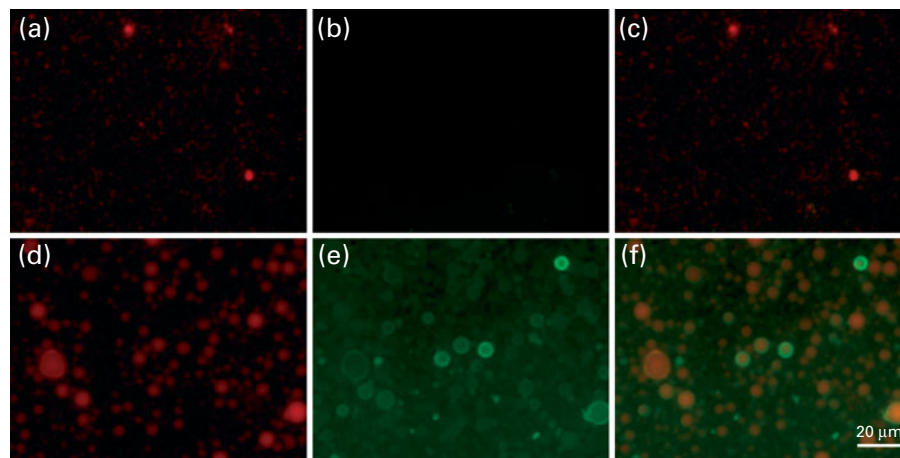


Fig. 2. Images of conventional epifluorescence microscopy with Nile Red-stained fat core (a, d), Annexin-V Alexa Fluor® 488-stained PL (b, e) and composed image of both (c, f) the Control and Concept IMF, respectively.

Table 3. Lipid droplet size characteristics of the Control infant milk formula (IMF) and Concept IMF derived from the particle size distribution analysis

	Control IMF	Concept IMF
$D(0.1)^*$ (μm)	0.183	0.407
$D(0.5)^*$ (μm)	0.349	4.951
$D(0.9)^*$ (μm)	0.855	11.67
$D(1.0)^*$ (μm)	1.633	23.24
$D[3, 2]^\dagger$ (μm)	0.285	1.337
$D[4, 3]^\ddagger$ (μm)	0.468	6.000
$< 1 \mu\text{m}^\S$ (%)	94.34	12.59
$2-12 \mu\text{m}^\parallel$ (%)	0.000	70.30
Mode ¶ (μm)	0.439	6.500

* 10, 50, 90 and 100 % particle volume percentiles.

† Surface-weighted average diameter.

‡ Volume-weighted average diameter.

§ Sum volume of particles $< 1.000 \mu\text{m}$ (% total volume of all particles).

$^\parallel$ Sum volume of particles of $2-12 \mu\text{m}$ (% total volume of all particles).

¶ Peak diameter of particle size distribution.

in the Concept IMF mice compared with the Control IMF mice (Fig. 5).

Plasma metabolic profile. Although both the IMF groups had higher concentrations of TC, HDL-C and LDL-C compared with the non-challenged Reference group ($P < 0.05$), the Concept IMF mice had markedly lower concentrations compared with the Control IMF mice ($P < 0.05$). Experimental diets did not significantly affect any of the analysed parameters for glucose homeostasis (Table 5).

Effect of the Western-style diet challenge during adolescence and adulthood

To confirm that the mild WSD indeed represented a diet challenge that affects adult phenotype, we compared the Control IMF group fed the WSD in adulthood with a non-challenged Reference group that was switched to a standard rodent chow on PN42.

WSD intake enhanced BW gain (17%) in the Control IMF mice compared with the non-challenged Reference mice (Fig. 3; $P = 0.002$) from PN42 to 98. Lean body mass gain was similar in both the groups ($F(2,31) = 0.056$, $P = 0.956$), but the WSD challenge increased FM by 80% and %FM by 50%. The weights of *m. tibialis*, pancreas and brain were unaffected (Table 4), whereas average liver weight of the Control IMF mice was higher compared with the Reference mice ($F(2,31) = 5.656$, $P = 0.008$).

The WSD challenge resulted in more large adipocytes in the Control IMF group compared with the Reference group. Average EPI adipocyte volume was 94% increased as a result of the 56 d WSD challenge, whereas average ING adipocyte volume appeared to be only 17% higher following the WSD challenge. This difference did not reach significance (data not shown). The Control IMF and Reference groups had comparable adipocyte numbers in both the EPI and ING depots (data not shown).

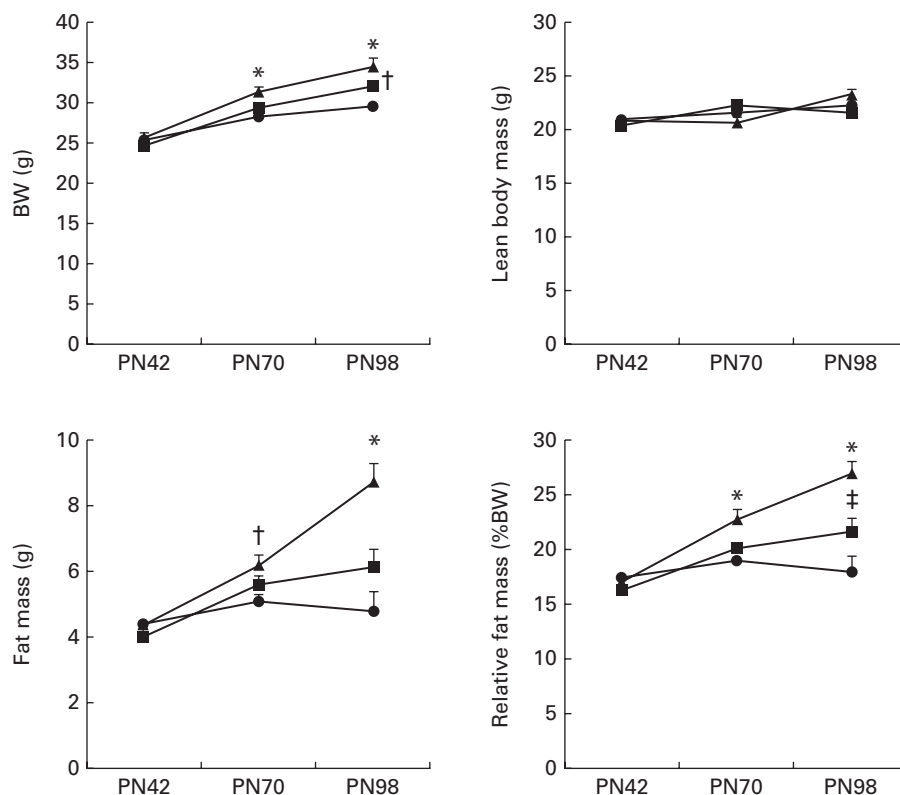


Fig. 3. Development of body composition ((a) body weight (BW), (b) lean body mass, (c) fat mass and (d) relative fat mass (%BW)) during the Western-style diet challenge of mice fed the early Control IMF (▲; n 11) or Concept IMF (■; n 11) and the Reference group (●; n 12) fed the American Institute of Nutrition-93M diet from postnatal day (PN)42 to 98. Values are means, with their standard errors represented by vertical bars. * Mean value was significantly different from that of the Reference and Concept IMF groups ($P < 0.05$). † Mean value was significantly different from that of the Reference group ($P < 0.05$). ‡ Mean value was significantly different from that of the Reference and Control IMF groups ($P < 0.05$).

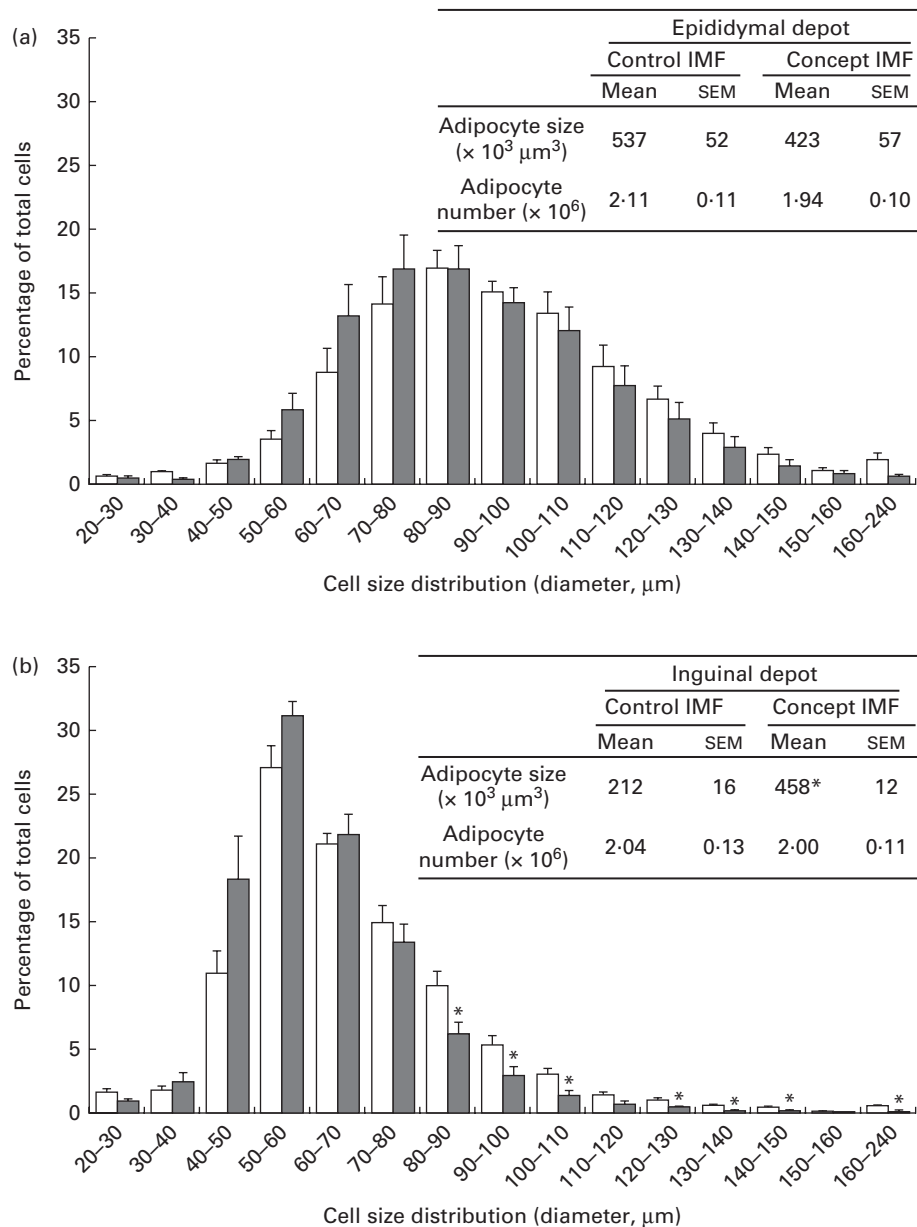


Fig. 4. Frequency distribution of adipocyte size, average cell size and number of epididymal (a) and inguinal (b) white adipose tissue depots on postnatal day (PN)98 in the Control infant milk formula (IMF) (□) and Concept IMF (■) groups. Values are means (n 9–12 mice per group), with their standard errors represented by vertical bars. * Mean value was significantly different from that of the Control IMF group ($P < 0.05$).

Fasting plasma TC ($F(2,27) = 17.352$, $P < 0.001$), HDL-C ($F(2,27) = 19.686$, $P = 0.049$) and LDL-C ($F(2,27) = 18.424$, $P < 0.001$) concentrations were increased in the WSD-challenged Control IMF mice compared with the non-challenged Reference group. Fasting plasma TAG, glucose, insulin and homeostasis model assessment of insulin resistance concentrations were not affected on PN98 due to the WSD challenge; the concentrations of the Control IMF and Reference mice were not significantly different (Table 5).

Discussion

The present study demonstrated that the physical structure of dietary lipids programmes body composition development

of pre-weaning mice towards reduced body fat accumulation in adulthood. These findings are in line with our previous study⁽²²⁾, despite differences in FA composition, indicating that the physical lipid structure indeed is the main determinant of the programming effect. The programming of adult FM by the Concept IMF was not mediated by a reduction in the number of adipocytes, but rather by a reduction of adipocyte size, indicating reduced lipid storage. The phenotype coincided with a reduced expression of transcription factors that promote lipogenesis in WAT.

Reduced adult fat mass and lipogenic gene expression

The development of human WAT starts in the third trimester of gestation and continues in postnatal life through the

Table 4. Average weight of white adipose tissue (WAT) depots and organs on postnatal day (PN)98 during the Western-style diet challenge of mice fed the Control infant milk formula (IMF) or Concept IMF and the Reference group fed the American Institute of Nutrition-93M diet from PN42 to 98

(Mean values with their standard errors, n 11–12)

	Reference		Control IMF		Concept IMF	
	Mean	SEM	Mean	SEM	Mean	SEM
WAT						
Epididymal (mg)	516.8	39.9	1138.0*	99.7	827.2*†	106.5
Perirenal (mg)	26.2	2.7	67.4*	8.3	43.8*†	5.5
Retroperitoneal (mg)	137.5	10.4	299.8*	26.3	215.2*†	29.5
Inguinal (mg)	374.8	32.0	542.8*	40.3	401.7†	28.4
Liver (g)	1.41	0.0	1.69*	0.1	1.57	0.1
Pancreas (mg)	159.3	8.0	167.5	7.8	172.1	9.5
<i>M. tibialis</i> (mg)	50.4	2.3	49.2	0.8	50.9	0.8
Brain (mg)	425.8	7.6	413.0	7.1	422.2	6.0

* Mean value was significantly different from that of the Reference group ($P < 0.05$).

† Mean value was significantly different from that of the Control IMF group ($P < 0.05$).

proliferation of preadipocytes and subsequent differentiation, resulting in mature adipocytes storing energy in the form of lipids⁽¹⁾. The amount of total body FM is determined by adipocyte size and by the number of differentiated adipocytes in WAT^(2,28). Feeding the Concept IMF for a relatively short period in early life partially prevented excess body fat accumulation upon a WSD challenge in later life. The Concept IMF mice exhibited a reduced storage of TAG compared with the Control IMF mice. The reduced gene expression of *leptin* in the Concept IMF mice compared with the Control IMF mice reflects both reduced FM and reduced adipocyte size⁽²⁹⁾. The reduced lipid accumulation upon the WSD challenge in the WAT of adult mice in the Concept IMF group was associated with reduced expression of *PPARγ* and *RXRα* mRNA in WAT, suggesting reduced FA uptake and/or *de novo* lipogenesis.

The expression of another key regulator of (*de novo*) FA and TAG synthesis, SREBP1c, was not affected by the postnatal diet. Possibly, the Concept IMF did not have a sustained impact on the expression of SREBP1c in WAT. Alternatively, the expression of SREBP1c is up-regulated by insulin, resulting in increased FA synthesis and reduced FA oxidation after feeding⁽³⁰⁾. In the present study, the expression of SREBP1c was measured after 4 h of fasting, which may have blunted possible differences between the groups in fed state⁽³¹⁾. Additionally, as SREBP1c is expressed in both WAT and liver, the Concept IMF may have exerted organ-⁽³²⁾ and WAT depot-specific⁽³³⁾ effects. Indeed, it is suggested that SREBP1c in WAT regulates the expression of genes involved in cholesterol metabolism rather than that of genes involved in FA and TAG synthesis⁽³⁴⁾.

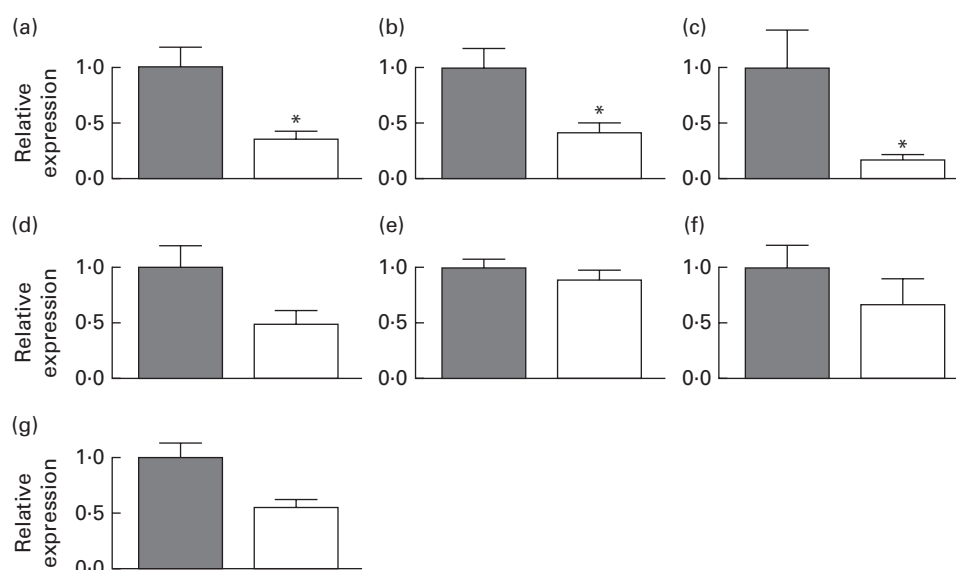


Fig. 5. Relative mRNA expression of (a) *PPARγ*, (b) *RXRα*, (c) *CEBPα*, (d) *CEBPδ*, (e) *CEBPβ*, (f) *SREBP1C* and (g) *leptin* in the epididymal white adipose tissue depot of the Control infant milk formula (IMF) (■) and Concept IMF (□) groups. Values are means (n 12 mice per group), with their standard errors represented by vertical bars. * Mean value was significantly different from that of the Control IMF group ($P < 0.05$).

Table 5. Average fasting plasma parameters on postnatal day (PN)98 during the Western-style diet challenge of mice fed the Control infant milk formula (IMF) or Concept IMF and the Reference group fed the American Institute of Nutrition-93M diet from PN42 to 98

(Mean values with their standard errors, n 2–4 mice per group on PN42 and n 8–11 mice per group on PN98)

	Reference		Control IMF		Concept IMF	
	Mean	SEM	Mean	SEM	Mean	SEM
PN42						
TAG (mmol/l)	1.66	0.1	1.44	0.1	1.85	0.1
TC (mmol/l)	3.48	0.1	3.58	0.2	3.18	0.1
HDL-C (mmol/l)	1.84	0.0	1.88	0.2	1.63	0.1
LDL-C (mmol/l)	0.89	0.1	1.04	0.1	0.71	0.1
VLDL-C (mmol/l)	0.76	0.1	0.66	0.0	0.84	0.1
PN98						
TAG (mmol/l)	0.97	0.1	1.00	0.1	0.82	0.0
TC (mmol/l)	2.98	0.2	4.40†	0.2	3.79*†	0.1
HDL-C (mmol/l)	1.76	0.1	2.63†	0.1	2.32*†	0.1
LDL-C (mmol/l)	0.77	0.1	1.31†	0.1	1.09*†	0.1
VLDL-C (mmol/l)	0.44	0.1	0.45	0.0	0.37	0.0
Glucose (mmol/l)	13.7	1.1	12.0	1.0	13.2	0.7
Insulin (pmol/l)	90.6	24.8	109.4	29.9	70.1	7.4
HOMA-IR ((mmol/l × pmol/l)/22.5)	42.0	12.6	56.8	14.3	42.0	5.8

TC, total cholesterol; HDL-C, HDL-cholesterol; LDL-C, LDL-cholesterol; VLDL-C, VLDL-cholesterol; HOMA-IR, homeostasis model assessment of insulin resistance.

* Mean value was significantly different from that of the Control IMF group ($P < 0.05$).

† Mean value was significantly different from that of the Reference group ($P < 0.05$).

Reduced adult fat mass reduced adipocyte size, but not adipocyte number

The Concept IMF mice had approximately 30% less total FM after the WSD challenge compared with the Control IMF mice, which was reflected in all the four WAT depots measured. However, the programming of adipocyte size by the Concept IMF appeared to be more pronounced in subcutaneous ING depots than in abdominal EPI depots, despite the comparable depot weight reduction. This difference could be explained by the major difference in EPI size distribution in the size category with the largest cells (150–240 μ m). These cells represent a small proportion of the total cell population, but store a relatively large amount of lipids compared with the smaller adipocytes. Thus, large adipocytes contribute significantly to the average adipocyte volume and storage capacity. It is known that there are WAT depot-specific differences in the responsiveness of adipocytes to nutrients, growth patterns, and metabolic and endocrine activities^(35–39). For instance, WAT expansion in ageing rats was associated with hypertrophic growth in visceral depots and hyperplastic growth in subcutaneous depots⁽³⁵⁾. Additionally, adipocyte size, number and gene expression have been shown to differ considerably between WAT depots in adult rodents^(40,41), which is in accordance with findings from human studies^(42–44). Several cohort studies have shown that fetal and postnatal growth velocity differentially affects human body fat distribution, with specifically rapid postnatal weight gain being strongly associated with visceral adiposity^(45–48). This suggests that the critical window for WAT development may be different for visceral and subcutaneous depots. Indeed, data from a Dutch infant cohort demonstrated that the amount of visceral body fat increased between 12 and 24 months of age, in contrast to the amount

of subcutaneous fat, which remained constant over this period⁽⁴⁹⁾. These age-dependent growth differences might also appear in rodents, as DiGirolamo *et al.*⁽¹⁰⁾ demonstrated distinct adipose tissue growth patterns among fat depots at different stages of life in rats. Additionally, both early-life overnutrition⁽⁵⁰⁾ and undernutrition⁽⁵¹⁾ can modulate these differential depot growth patterns. Unfortunately, we were unable to investigate adipocyte size distribution and number in visceral depots other than EPI WAT due to the small fat depot size, providing insufficient material for analysis. Therefore, the present study could not provide conclusive evidence for postnatal nutritional programming of adipocyte number by the Concept IMF. One could hypothesise that the nutritional intervention was implemented after a critical window for adipogenesis. Earlier dietary exposure to exclude this possibility would require an artificially fed animal model. However, early-life handling is known to induce stress as a confounding factor since it influences later-life body composition and fat distribution^(52,53). Alternatively, the absence of a sustained effect on adipocyte number could be related to the fact that both IMF-containing early diets meet the requirements for normal growth and development. Therefore, there would be no need for the mice to utilise the plasticity of WAT to increase adipocyte number in response to this nutritional stimulus. It has been hypothesised by Spalding *et al.*⁽³⁾ that increased adipocyte number in early life is a major risk factor for adult obesity. However, it is still unclear whether this is a causal relationship; it is unknown whether increased adipocyte number itself predisposes to adult obesity or whether both increased adipocyte number and adult obesity are caused by, for instance, excessive energy intake during childhood through a different mechanism altogether. In accordance with this hypothesis, early postnatal overnutrition in rats has been found to

specifically increase adipocyte number and thereby increase adult FM, but it did not affect adipocyte size⁽⁵⁴⁾. Additionally, hyperplasia and cell turnover capacity are positively correlated with insulin sensitivity independent of BW, whereas hypertrophy is negatively correlated with insulin sensitivity⁽⁵⁵⁾. Our data clearly indicate that early diets can induce programming activity that is independent of hyperplastic growth, but seems to affect the capacity of adipocytes to enlarge upon a WSD challenge.

As stated previously, critical periods for the hyperplastic growth of WAT encompass early postnatal life and adolescence^(2,56); cell number remains constant in healthy non-obese adults⁽³⁾. However, in case of sustained excessive energy intake in obesity, adult WAT is still capable of adjusting by both hypertrophic and hyperplastic growth⁽⁵⁷⁾. In the present study, the moderate WSD significantly increased adipocyte size compared with a standard chow, but was not such a severe challenge as to promote hyperplasia in adolescence and adulthood. We cannot exclude that hyperplasia would have occurred if a more severe dietary challenge was used. Thus, presently we cannot rule out the potential programming effects of the postnatal diet on hyperplastic WAT growth.

Programming of adult fat mass and metabolic homeostasis

We determined to what extent the observed differences in adult body composition and FM corresponded with other parameters of lipid homeostasis, such as plasma lipids. Plasma TC, HDL-C and LDL-C concentrations were reduced in the Concept IMF mice compared with the Control IMF mice. Liver weight of the Control IMF mice was greater than that of the non-challenged Reference mice, whereas liver weight of the Concept IMF mice did not differ significantly from that of both the groups. This may suggest an improved metabolic profile and perhaps the absence of ectopic lipid accumulation due to the Concept IMF. Since plasma TAG, glucose and insulin concentrations were not significantly perturbed by the moderate WSD challenge compared with a normal rodent chow, the limited improvement in adult glucose homeostasis and TAG concentrations of the Concept IMF-fed mice was to be expected.

Effect of milk lipid physical structure: potential mechanisms

Previous studies comparing the digestion of raw cows' milk with that of processed milk with different milk lipid droplet sizes and surface compositions have reported differences in gastric emptying, lipid digestion and β -oxidation rate^(58–60). The results suggest differences in the utilisation of these lipids for either β -oxidation or lipogenesis. In turn, this could underlie the early programming of lipid metabolism resulting in differences in the metabolic capacity of adipocytes.

The sustained effects of the Concept IMF on body composition and metabolic profile cannot simply be explained by reduced food intake. Although individual food intake during exposure to early diets could not be measured since it was provided as dough to the entire litters until weaning, BW and

body composition were similar between the groups on PN42. Moreover, daily food intake was comparable between the groups during the WSD challenge. Possibly, the Concept IMF-fed mice utilise energy from the WSD differently by storing less energy as lipids in adipose tissue, suggesting a higher BMR. However, this information is currently not available. Future studies that may help us gain more insights into energy expenditure are pivotal. Also, further studies addressing which specific aspects of the Concept IMF, i.e. fat droplet size, PL addition and/or coating, may contribute to the observed long-term effects are ongoing.

Conclusion

In conclusion, early exposure to the Concept IMF resulted in a diet-resistant phenotype in adult life. The Concept IMF prevents excessive hypertrophic growth of WAT in adulthood despite unaltered adipogenesis. Our data support the hypothesis that early Concept IMF exposure programmes metabolic responses through altered gene expression of factors regulating adipose tissue lipogenesis. The present results provide support for the emerging notion that dietary lipid structure should be considered a key component of dietary lipid quality in early life and is a determinant in later-life obesity risk.

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